Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claim 1. (Original): A compound of formula I

wherein

R_a is H; C₁₋₄alkyl; or C₁₋₄alkyl substituted by OH, NH₂, NHC₁₋₄alkyl or N(di-C₁₋₄alkyl)₂;

R_b is H; halogen; C₁₋₆alkyl; or C₁₋₆alkoxy, and

R is a radical of formula (a) or (b)

$$R_2$$
 R_4
 R_4
 R_3
(a)
(b)

wherein

each of R_1 and R_3 is a heterocyclic residue; or a radical of formula α

$$-X-R_c-Y$$
 (α)

wherein X is a direct bond, O, S or NR₁₁ wherein R₁₁ is H or C₁₋₄alkyl,

 R_c is C_{1-4} alkylene or C_{1-4} alkylene wherein one CH_2 is replaced by CR_xR_y wherein one of R_x and R_y is H and the other is CH_3 , each of R_x and R_y is CH_3 or R_x and R_y form together – CH_2 - CH_2 -,

Y is bound to the terminal carbon atom and is selected from OH, $-NR_{12}R_{13}$ wherein each of R_{12} and R_{13} , independently, is H, C_{3-6} cycloalkyl, C_{3-6} cycloalkyl- C_{1-4} alkyl, aryl, aryl- C_{1-4} alkyl, heteroaryl- C_{1-4} alkyl, C_{2-6} alkenyl or C_{1-4} alkyl optionally substituted on the terminal carbon atom by OH, halogen, C_{1-4} alkoxy or $-NR_{14}R_{15}$ wherein each of R_{14} and R_{15} , independently, is H, C_{1-4} alkyl, C_{3-6} cycloalkyl, C_{3-6} cycloalkyl- C_{1-4} alkyl, aryl- C_{1-4} alkyl, or R_{12} and R_{13} form together with the nitrogen atom to which they are bound a heterocyclic residue; and

each of R₂ and R₄, independently, is H; halogen; C₁₋₄alkyl; C₁₋₄alkoxy; CF₃; nitrile; nitro or amino,

or a salt thereof.

Claim 2. (Original): A compound according to claim 1 wherein R_a is H, methyl, ethyl, or isopropyl, or a salt thereof.

Claim 3. (Currently amended): A compound according to claim 1–or 2 wherein R_b is H, Cl, methyl or ethyl, or a salt thereof.

Claim 4. (Currently amended): A compound according to any one of claims 1-to 3 wherein R_1 is a heterocyclic residue, e.g. a piperazinyl, optionally substituted on a ring nitrogen or on a ring carbon, e.g. 4-methyl-piperazin-1-yl, or 4,7-diaza-spiro[2.5]oct-7-yl; or a radical of formula (α) wherein X is a direct bond, R_c is CH_2 and Y is $-NR_{12}R_{13}$ wherein each of R_{12} and R_{13} , independently, is H, C_{3-6} cycloalkyl- C_{1-4} alkyl, C_{2-6} alkenyl or C_{1-4} alkyl optionally substituted on the terminal carbon atom by OH, halogen, C_{1-4} alkoxy or $-NR_{14}R_{15}$ wherein each of R_{14} and R_{15} , independently, is H or C_{1-4} alkyl; or R_{12} and R_{13} form together with the nitrogen atom to which they are bound a heterocyclic residue e.g. a piperazinyl, or a salt thereof.

Claim 5. (Currently amended): A compound according to any one of claims 1 to 4 wherein R_2 and/or R_4 is H; Cl, F; CF_3 ; nitrile; nitro or amino, or a salt thereof.

Claim 6. (Original): A process for the preparation of a compound of formula I according to claim 1, which process comprises reacting a compound of formula II

wherein R_a and R_b are as defined in claim 1,

with a compound of formula III

$$R - CH_2 - CO - NH_2$$
 (III)

wherein R is as defined in claim 1,

and, where required, converting the resulting compound of formula I obtained in free form to a salt form or vice versa, as appropriate.

Claim 7. (Currently amended): A compound of formula I according to any one of claims 1 to 5, in free form or in a pharmaceutically acceptable salt form for use as a pharmaceutical.

Claim 8. (Currently amended): A pharmaceutical composition comprising a compound of formula I according to any one of claims 1-to-5, in free form or in a pharmaceutically acceptable salt form, in association with a pharmaceutically acceptable diluent or carrier therefor.

Claim 9. (Currently amended): A compound of formula I according to any one of claims 1 to 5 or a pharmaceutically acceptable salt thereof for use in the preparation of a pharmaceutical composition for use in the treatment or prevention of disorders or diseases mediated by T lymphocytes and/or PKC or GSK-3β.

Claim 10. (Currently amended): A method for preventing or treating disorders or diseases mediated by T lymphocytes and/or PKC or GSK-3β in a subject in need of such treatment, which method comprises administering to said subject an effective amount of a compound of formula I according to any one of claims 1 to 5 or a pharmaceutically acceptable salt thereof.